Study of Li$^+$ Ion-exchanger Mg$_2$TiO$_4$

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Abstract. Spinel-type metal oxides, magnesium-titanium oxide (Mg$_2$TiO$_4$), was prepared by a solid state reaction crystallization method. The spinel construction of Mg$_2$TiO$_4$ was conformed by XRD. The results showed that the Li$^+$ extraction/insertion be progressed mainly by an ion-exchange mechanism. The acid treated samples had an ion exchange capacity of 13.9mmol/g for Li$^+$.

Introduction

The inorganic ion-exchange preparation has the advantage of thermo-stability and radiation resistance, synthesis simple and good selectivity etc. In addition, it appears the fine speciality in dealing with nuclear waste, gathering and separating of metal ions and chromatogram analysis[1-2]. In this paper, inorganic ion exchanger (Mg$_2$TiO$_4$) with a spinel structure ion memory was synthesized by a solid state reaction crystallization method., which is different the reported in the literatures[3-5], and its ion-exchange properties were studied.

Experimental Section

Reagent and Instruments. MgO and TiO$_2$ were all analytical reagents; pure ethanol; D/max-A type X-ray diffraction instrument; Dx-170 type ion chromatogram instrument; XQM planetary ball mill; AA-670 atom absorption spectrum instrument; tubular-furnace.

Synthesis and Identifiable of Mg$_2$TiO$_4$. The pure ethanol was dropped into a XQM planetary ball mill mixed powder of MgO and TiO$_2$ with a Mg/Ti mole ration of 2:1 at the condition of constant rate churning. After 8 hours, the mixture was mixed completely. After mixing fully, the mixture was pressed to tablet by tablet press machine. Then the tablet was heat-treated for 4.5h at 900°C to obtain the Mg-Ti metal compound, the sample was designed as MgTi-900, whose theoretical formula was Mg$_2$TiO$_4$. Then it was analysed of x-ray diffraction and compared to literature[3-4].

Composition analysis: A 0.2g portion of sample was dissolved with acid. The Mg and Ti contents were determined by atomic absorption spectrometry.

The Cation Extraction of Mg-Ti metal compound and Acid Modification. Four 0.200g portions of sample (MgTi-900 compound) were immersed in a HNO$_3$ solution (50ml) of 0.01M, 0.1M, 1M and 10M respectively with shaking in constant temperature water at 25°C. After 3 days, take the supernatant solution to determine the cation concentration, test its acid proof ability and the extraction ration of Ti$^{4+}$.

A 5g portion of sample (MgTi-900) was immersed in a 1M HNO$_3$ solution (500mL) with intermittent shaking in constant temperature water at 25°C. After 7 days, remove the supernatant solution and add new HNO$_3$ solution. Repeating that for twice, then the initial sample was transformed to H-type sample, washed with water and air-dried. The sample obtained by thermal crystallized at 900°C and acid modified was designated as MgTi-900 (H).

Saturation Capacity of Exchange. Weigh five 0.5g portions of MgTi-900 (H), then each portion was immersed in a 0.1M solution (10mL), containing Li$^+$, Na$^+$, K$^+$, Rb$^+$ and Cs$^+$ respectively, diluted to 100mL, shaken in constant temperature water at 25°C. After saturation exchanging (namely, after 10 days by literature[3-4]) the solutions were filtered by subminiature aperture sieve, and the cation concentration was determined. At the same time, do vacant experiment. Last, the
inorganic exchanger saturation capacity of exchange for alkali-metal-ions obtained by decrease quantity.

**Distribution Coefficient (Kd).** After weighing four 0.100g portions of MgTi-900 (H), each portion of sample was immersed in a 0.05M mixed solution (0.200mL) containing Li⁺, Na⁺, K⁺, Rb⁺ and Cs⁺ (Cl⁻/OH⁻ ratios are different in each solution, C(Cl⁻)+C(OH⁻)=0.1M, C=Li⁺, Na⁺, K⁺, Rb⁺ and Cs⁺). The alkali-metals ions total concentration all was $1.0 \times 10^{-3}$M by adding 9mL distilled water. After the samples were shaken for 7 days in constant temperature water at 25°C and were filtered, cation concentrations in each samples were obtained.

**Results and Discussion**

**Compound and Appraisal of Mg₂TiO₄.** The X-ray diffraction pattern of compound metal oxide (Mg₂TiO₄), crystallized was shown in figure 1. The structure of compound metal oxide Mg₂TiO₄ crystallized at 900°C was much perfect.

We know from chemical analysis, the composition of MgTi-900 is Mg₁.₉₇Ti₀.₉₉O₃.₉₆, whose chemical component is basically corresponded with the composition of spinel-type metal oxides.

![Fig.1 The powder’s X-ray figures of Mg₁.₉₇Ti₀.₉₉O₃.₉₆ crystal](image)

**Cation Extraction of Metal Compound Mg-Ti and Acid Modification.** The extraction ration of Mg²⁺ and Ti⁴⁺ from MgTi-900 in different concentration HNO₃ solution is shown in Figure2. We know from Figure2, the extractabilities of Mg²⁺ are 35%~78% and Ti⁴⁺ are 2.4%~8.0%. Those indicate that the extractabilities of Mg²⁺ are higher than those of Ti⁴⁺ when exchanger was immersed in 1M acid solution, corresponding with the exchanger condition was better.(1 N, Mg²⁺73%, Ti⁴⁺6.1%)

X-ray diffraction of MgTi-900 (H), which is the acid modification product, is shown in Figure1. As shown, the structure of MgTi-900(H) is nearly constant, which is spinel oxide type too. It indicate that the exchanger is steady. The analysis indicate the composition of MgTi-900 (H) was H₂.₈₇Mg₁.₃₃Ti₀.₉₃O₃.₆₄, whose component of 73% Mg²⁺ transformed to H⁺ compared with the composition Mg₁.₉₇Ti₀.₉₉O₃.₉₆ before acid-treated. Then the specific Mg²⁺ of exchanger were extracted fulfill basically and remained the H-type identified with initial type.
Saturation Capacity of Exchange. The relation between radius and saturated ion exchange capacity of MgTi-900 (H) for alkali was shown in figure 3. Known from figure 3, the capacity of exchange for Li⁺ was much higher than those for other alkali ions. The capacity for Li⁺ is 13.9 mmol·g⁻¹. It proved that the ion exchange synthesized has higher capacity of exchange, and better remembering of exchange for Li⁺. The effect factors of saturation capacity of exchange of MgTi-900 (H) are: 1) The Li⁺ in exchange solution must be removed previously, because Li⁺ exchanged with exchanger vacancy site when existing too much Li⁺; 2) The experimental results shown that the exchange capacity of ion exchanger for Li⁺ is much higher than those for other alkali ions in thin solution, which indicate that the ion-exchange reaction is carried out between and bare ions; 3) At the time of exchange, a Li⁺ was replaced by one H⁺. Li⁺ not only entered the vacancy site but also exchanged with the H⁺ of surface. Therefore, MgMnTi-900 (H) has a higher exchange capacity for Li⁺.
**Distribution coefficient (Kd).** Kd values can be the token of exchange selectivity of MgTi-900 (H) for correlate ions. Shown in figure 4, Kd values of MgTi-900 (H) for alkali ions are larger and larger with an increase pH over the pH region studied. The selectivity sequence of MgTi-900 for alkali metal ions as follows:

\[ \text{Li}^+ > \text{Cs}^+ > \text{Rb}^+ > \text{K}^+ > \text{Na}^+ \] (1)

It indicates that MgTi-900 (H) has a better ion selectivity for Li\(^+\). Ion-exchange reaction is reversible reaction. The reaction of H\(^+\) in ion-exchanger with alkali metal ions in solution as follows (example for Li\(^+\)):

\[ E - 2H + 2\text{Li}^+ \Leftrightarrow E - 2\text{Li} + 2H^+ \] (2)

![Fig.4 Distribution coefficient of MgTi-900(H) for alkali ions](image)

**Conclusions**

The Mg\(_2\)TiO\(_4\) of spinel-type metal oxide show a capacity extraction/insertion of Li\(^+\) in the aqueous phase, by X-ray, saturation capacity of exchange and Kd measurement, etc. It has high selectivity to Li\(^+\). The Li\(^+\)-extracted samples show a high selectivity and a large capacity for Li\(^+\) among alkali metal ions.

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**References**

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